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300 Mcs by the coaxial-line method. Other specific properties of BiFeO₃ in addition to the large internal fields and large spontaneous polarization, were a high Curie temperature (850C), and the absence of nonlinear properties at room temperature. An analysis of all the published data and the present results shows that in spite of the fact that the spontaneous electric polarization of BiFeO₃ is very high, so that no dielectric hysteresis loops could be obtained, this substance is ferroelectric. Various reasons for this conclusion are discussed. An analysis of the magnetic properties below the Neel temperature (370C) also shows that BiFeO₃ has compensated ferromagnetism in addition to ferroelectricity. Orig. art. has:

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THE COMPANY HER SECRETARY TO CARE DATA AND SECRETARIZED BY SECRETARIZED AND SECRETARIZED BY SE EWT(m)/T/EWP(t)/ETI IJP(c) SOURCE CODE: UR/0181/66/008/009/2594/2597 44 L 01052-67 В AP6030956 AUTHOR: Kashlinskiy, A. I.; Chechernikov, V. I.; Venevtsev, Yu. N. ACC NR: ORG: Moscow State University im. M. V. Lomonosov (Moskovskiy gosudarstvennyy universitet) TITLE: Investigation of electron resonance and magnetic properties in solid solutions of the system SOURCE: Fizika tverdogo tela, v. 8, no. 9, 1966, 2594-2597 TOPIC TAGS: electron spin resonance, electron spectrum, EPR spectrum, solid solution, bismuth ferrate, barium titanate ABSTRACT: The spectra of electronic resonance in solid solutions of the system bismuth ferrate barium titanate have been investigated. The clearly defined anomalies in the spectra are determined, corresponding to the dielectric and magnetic transitions in solid solutions in conformity with tetragonal and rhombohedral modifications. The data on changes in the EPR spectra are analyzed in relation to the properties of solid solutions under study. Orig. art. has: 2 figures. [Based on authors' abstract; SUB CODE: 20/ SUBM DATE: 14Jan66/ ORIG REF: 009/ Card 1/1 awm

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CIA-RDP86-00513R001859410008-5

10001-112001 v. Venevtsey, Yu. N., . 1540. A. = . ORG: none TITLE: Mossbauer effect in the ferroelectric Po(Fe1/2No1/2)03 SOURCE: Zhurnal eksperimental noy i teoreticheskoy fiziki. Pis'ma v redaktsiyu. Prilozheniye, v. 3, no. 5, 1966, 212-216 TOPIC TAGS: ferroelectric material, Messbauer spectrum, miltiplet splitting, critical point, phase transition, Curie point, electron spin ABSTRACT: The purpose of the investigation was to check on the presence of a minimum of the probability of the Mossbauer effect on Sn119 in the investigated compound, similar to that observed for Ba(TiSh)O3 (with different Ti/Sh ratios) by V. A. Bokov et al. (FTT v. 7, 1886, 1965 and elsewhere). It was also desired to check on other singularities in the behavior of the quadrupole splitting and of the position of the symmetry center of the Mossbauer spectrum observed near the temperature $T_{\rm C}$ of the ferroelectric phase transition. To this end, the authors investigated the variation of the parameter of the Mossbauer absorption spectrum of Fe^{57} nuclei of the ferroelectric in question at the phase transition temperature ($F_c = 11^h C$). The absorbers were made by the usual ceramic technology, using $Fe_2^{57}O_3$ (Fe^{57}). The source was Co57 in stainless steel. The apparatus for the Mossbauer spectra is described by the authors elsewhere (PTE No. 4, 43, 1964). The results confirm the existence of the

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医全球性动物 医结节性结合性 医生物性结合 医神经神经病 化多种性多种性多种性多种性多种性 L 24372-66 ACC NR: APEO10437 singularities in the vicinity of the Curie point $T_{\rm C} = 214{\rm C}$ and a minimum in the Mossbauer-effect probability. These singularities are apparently connected with the fact that an anomalous decrease in the frequency of one of the transverse optical branches of the lattice takes place on approaching the ferroelectric transition point in crystals with perceskite structure. The decrease in the quadrupole splitting with increase of temperature to To is connected with a decrease in the spontaneous polarization. The asymmetry of the quadrupole-splitting line, which has a minimum near $T_{\rm C}$ The Wishauer effect probability investigation of the toformation on the dynamics of the second of the constant structure ferroelectric transition. The authors had a few churreyev and Man's in the ferroelectric transition. The authors had a few control of the few controls and the few controls are sent to the few controls and the few controls are sent to the few controls and the few controls are sent to the few controls and the few controls are sent to the few controls are sent to the few controls and the few controls are sent to the ing are a wear meter. It is the transfer THE HEAVING

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SOURCE CODE: UR/0363/67/003/001/0268/0269

AUTHOR: Shapiro, Z. I.; Fedulov, S. A.; Venevtsev, Yu. II.

ORG: Physicochemical Institute im. L. Ya. Karpov (Fiziko-Ahimicheskiy institut)

TITIE: Determination of the Curie temperature of the ferroelectric LineO3

SOURCE: AN SSSR. Izvestiya. Neorganicheskiye materialy, v. 3, no. 1, 1967, 208-209

TOPIC TAGS: Curie point, lithium compound, niobate, ferroelectric crystal, dielectric constant

AESTRACT: In order to refine the Curie point of LiNbO3, temperature measurements of the dielectric constant were made on single crystals of both LiNbO3 and a solid solution of the composition Li(NbO.9TaO.1)O3. The Curie temperature, determined from dielectric constant maxima, was found to be 1210±10°C for LiNbO3 and 1120±10°C for the solid solution. In LiNbO3, the dielectric constant along the polar axis is much less than in the perpendicular direction, as in the case of barium tit mate single crystals. Some anemalies in the dielectric constant were found in the 600-950°C range. Thermographic measurements showed the melting point of LiNbO3 to be 1245±5°C. Of all known ferroelectrics, LiNbO3 has the highest Curie point. The data obtained on the Curie and melting points of LiNbO3 are of major importance for the preparation of single-domain single crystals with higher dielectric, optical and piezoelectric properties.

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Authors ar	re grateful to G	. P. Kuznots	ova, V. L. Fa	rshtendiker a	nd R. M. Tole	chinskaya
for assist	tance in the det	ermination o	f the melting	point, Orig	. art. hasi	l figure.
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ACC NR: AP6006823 SOURCE CODE: UR/0181/66/008/002/0416/0423

AUTHOR: Viskov, A. S.; Venevtsev, Yu. N.

ORG: Physicochemical Scientific Research Institute im. L. Ya. Karpov, Moscow (Nauchno-issledovatel'skiy fiziko-khimicheskiy institut)

TITLE: Calculating the gradients of the intracrystalline field in barium titanate ferroelectric and in model crystals based on this salt

SOURCE: Fizika tverdogo tela, v. 8, no. 2, 1966, 416-423

TOPIC TAGS: barium titanate, crystal theory, ferroelectric crystal

ABSTRACT: A method based on the ion model is proposed for calculating the gradients of the intracrystalline field VE which act on the ions in the tetragonal modification of ferroelectric barium titanate. The structural coefficients used in calculating the gradients were taken from experimental data in the literature. Curves lating the gradients were taken from experimental data in the literature. Curves are given showing the difference in lattice parameters (c-a) as a function of temperature. The effect which temperature has on spontaneous polarization, the intracrystalline field and the gradients of the field is analyzed for the same modifi-

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cation of barium titanate. The gradients are studied as a function of charge and polarizability of A and B cations in ABO3 crystals of the BaTiO3 type. The proposed method gives results which agree satisfactorily with the experimental data in the literature. The ion model is found to be highly effective for evaluating in the literature. The ion model is found to be highly effective for evaluating electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and gradients in ferroelectrics which are basically metal electric field intensities and

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TOMASHPOL'SKIY, Yu.Ya.; VENEVTSEV, Yu.N.

Perromagnetism in 'he system Pb₂CoWO₆ -- BaTiO₃. Fiz. tver.
(MIRA 18:11)
tela 7 no.10:3126-3128 0 '65.

1. Nauchno-issledovatel'skiy fiziko-khimicheskiy institut imeni
Karpova, Moskva.

RCGINSKAYA, Yu.Ya.; VEMENTORY, Yu.N.

Structure and reports properties of estimated in the system BiFeOg - InCr(3 Fiz. twer. talk y in addition if 165.

1. Nauchno-isoladovatelicity finite-directly institut iconi Karjova, Yoskva.

TOMASHPOLIUKIY, Yu.Ya.; VEHEVTSEV, Yu.N.; ANTONOV, G.N.

Ferromagnetic substances in the system PbFe2/3/1/303 Fb2YbNbC6. Zhur. eksp. i teor. fiz. 49 no.2:367-372 Ag *65. (MIRA 18:9)

1. Fiziko-khimicheskiy institut imeni Karpova.

VISKOV, A.S.; VENEVTSEV, Yu.N.; ZHDANOV, G.S.

New ferroelectric substances with the structure of perovskite and pyrochlore. Dokl. AN SSSR 162 no.2:323-325 My *65. (MIRA 18:5)

1. Fiziko-khimicheskiy institut im. L.Ya.Karpova. Submitted December 2, 1964.

KAPYSHEY, A.G.; VENEVISEV, Yu. U.

X-ray diffraction study of variations in the period of elementary cells of (Ba, Pb)TiO₃ solid solutions in the paraelectric region as dependent on the composition. Kristallografiia 8 no.2:269-270 (MIRA 17:8)

1. Fiziko-khimicheskiy institut imeni Karpova.

TOMASUROLISKIY, Yu.Ya.; VETEVICAY, Yu.W.

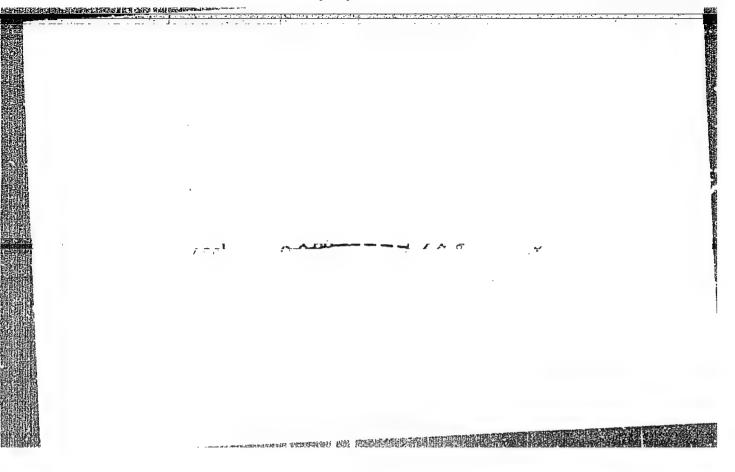
New lead-containing perceptites. Fin. terr. tola 7 no.2:517-520 (MURA 18:8)

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1. Mauchno-ideledevatel ship fiziko-khimicheskiy institut ineni Karpova, Moskva.

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ROGINSKAYA, Yu.Ye.; VENEVISEV, Yu.N.

Structure and dielectric properties of Pb2CcWC6. Kristellografiia 10 no.3:341-345 My-Je 165. (MIRA 18:7)

l. Fiziko-khimicheskiy institut imeni L.Ya. Karpova.

EMP'el'EPA's 1-2/EMP(+1/EMP(1)/EPA(W)-2/EMP(t)/EMP(b) LIP(c) 19 0070 65 010 005 064 1 0649 1 1,253-66 548, 736, 537, 246 ACC NR. AP5024548 AUTHOR: Viskov, A. S., Venevstev, Yu. N. Zhdanov, G. S. TITLE: Study of the structure and magnetic and electric properties of solid solutions in the system BiFeO₃ - "Sr(Sn₁/₃Mn₂/₃)O₃" SOURCE: Kristallografiya, v. 10, no. 5, 1965, 644-649 TOPIC TAGS: solid solution, Neel temperature, Curie point, antiferromagnetic material, spontaneous magnetization, bisruth compound, iron compound, strontium compound, tin compound, manganese compound 1) ABSTRACT: The samples were prepared by the usual ceramic process from Bi₂O₃, SnO₂, SrCO₃. Fe₂O₃, and MnO₂ (firing at 75 - 1100C at 4 - 6 hr and again at 820 - 1600C for 1-2 hr). After each firing, the phase composition of the samples was cheesed by x-ray diffraction. The latter showed that the system Rikings of Polyrages with the system Rikings of Polyrages of the state of the system and the polyrages of the state of the system and the s property of the spatting the spatting the contractions of their computitions and the consense sections exist in four modifications of the second of Neel temperatures is, of the virgous samples were determined in a 25 to 1977 pendence of spontaneous magnetization, the low the solid solid one are sender smarnetics with a weak ferromagnetism. It was found that in the system studied there is a Apple in the distance of

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solid solutions in which a magnetic and an electric dipose structure exist over a sole temperature range. No distinct correlation was noted between these structures, and no special effort is made to elucidate it. We thank Ya. Ye. Roginshy for variable suggestions and comments." Orig. art. has: 4 ligures.

ASSOCIATION: Fiziko-khimicheskiy institut im. L. Ya. Karpova (Physicochemical Institute)

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 Structure and min the system F 1022-1025 Je	agnetic properties Pb2CoWO6 - CdMnO3. 65.	of ferroelectric s Izv. AN SSSR. Ser.	fiz. 29 no.6: (MIRA 18:6)
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SHAPHer, Z.I.; VEIKLOV, S.A.; VENEVICET, TR.H.; RIGEROAN, L.G.

Study of the system LiTaC3 - Little 2. Lzv. An Shar. Ser. fiz.
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1. Vsesoyuznyy nauchno-issledov.tel*skiy institut khimicheskith reaktivov.

ROGINSKAYA, Yu.Ye.; VEHEVISEV, Yu.K.; VHDANOV, G.S.

New ferromagnetic substances. Thur. eksp. i teor. fir. 45 ix

1824-1232 My 165.

1. Nauchno-issledovatel'skiy fiziko-khimicheskoy institut insti
I. Ya. Karpova, Hoskva.

AUTHOR: Tomashpol'skiy, Yu. Ya.; Venevtsev, Yu. N.; Antonov, G. N.

TITLE: Ferroelectric-magnetic materials in the system PbFe2/3W1/3O3--Pb2YbNbO6

SOURCE: Zhurnal eksperimental noy i teoreticheskoy fiziki, v. 49, no. 2, 1765, 376-372

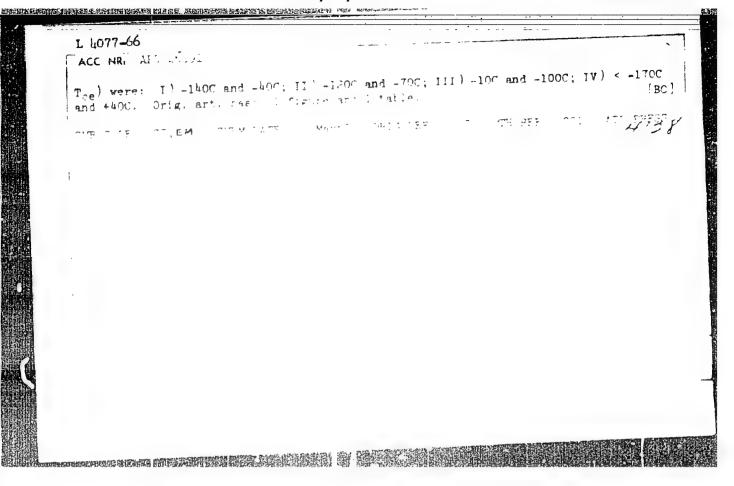
TOPIC TAGS: ferroelectric material, antiferromagnetic material, lead containing alloy, iron containing alloy, tungsten containing alloy, niobium containing alloy, solid solution

ABSTRACT: New ferroelectric-magnetic materials with perovskite structure were prepared in polycrystalline form by firing the oxides WO₃, Fe₂O₃, Nb₂O₅, and Yb₂O₅ and the carbonate PbCO₃ at 700--1000C for 1--7 hours. The techniques used for the crystal-structure measurements at high temperatures and for the phase measurements were described by the authors earlier (FTT v. 6, 298, 1964 and Zav. lab. no. 9, 1112, 1961). The dielectric measurements were made by a standard bridge method, and the magnetic measurements by the Faraday method. The tests showed that several solid solutions are formed in the PbFe_{2/3}W_{1/3}O₃--Pb₂YbNbO₆ system, some of which combine ferroelectric or antiferroelectric properties with ferrimagnetic ones over a relatively wide range of concentrations and temperatures. Some results were ob-

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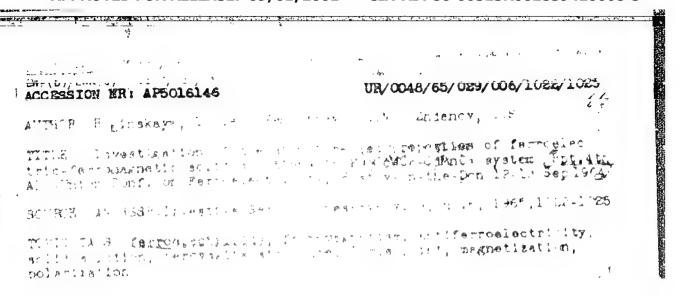
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	AUTHOR: Shvorneva, L. I.; Venevtsev, Yu. N.
	ORG: Physicochemical Institute im. L. Ya. Karpov (Fiziko-khimicheskiy institut)
	TITLE: Perovskites with ferroelectric properties
	SOURCE: Zhurnal eksperimental'noy i teoreticheskoy fiziki, v. 49, no. 4, 1965, 1038-1041
	TOPIC TAGE: ferroelectric material, ferroelectric property perovskite, inorganic synthesis
	ABSTRACT: In view of the important practical applications of ferroelectric compounds and of the necessity for a closer study of their properties, specimens of single-phase percyskites of the foll wing matter share test sections and the composition of the following matter sections and the composition of the following matter sections and the following matter sections are sections and the following matter sections and the following matter sections are sections and the following matter sections are sections and the following matter sections and the following matter sections are sections and the following matter
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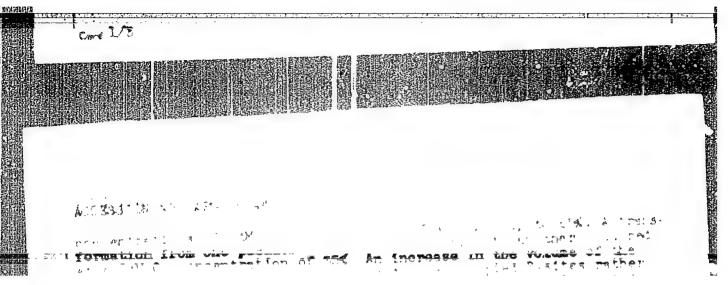


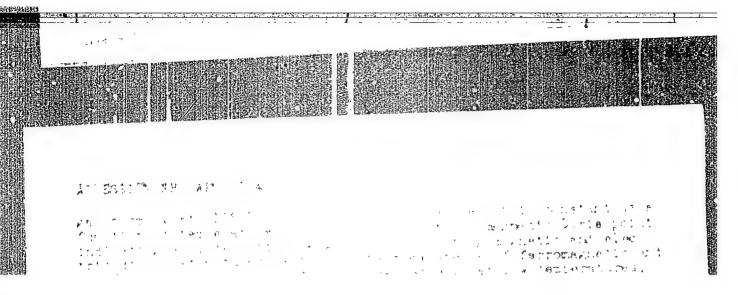
KAPYSHEV, A.G.; VENEVISEV, Yu.N.; SOLOV'YEV, S.F.; GORBUNOV, L.A.; ZHDANOV, G.S.

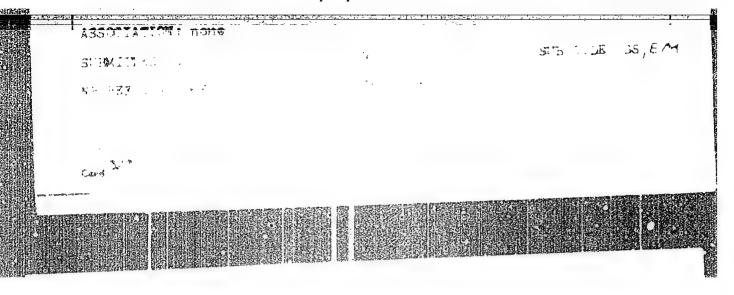
X-ray chambers for high-temperature studies. Zav. lab. 3C no.10: 1274-1276 *64. (MIRA 1834)

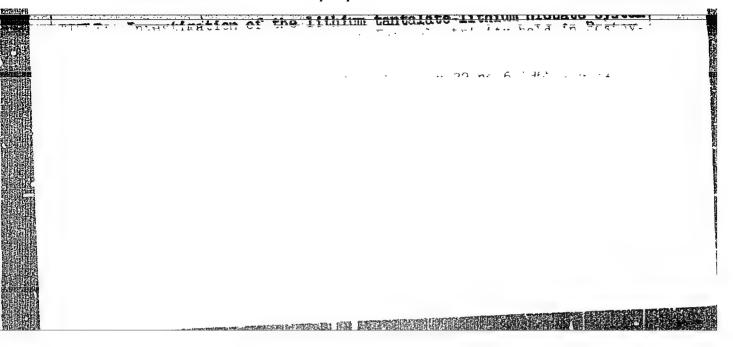
1. Nauchno-issledovatel skiy fiziko-khimicheskiy institut imeni Karpova.



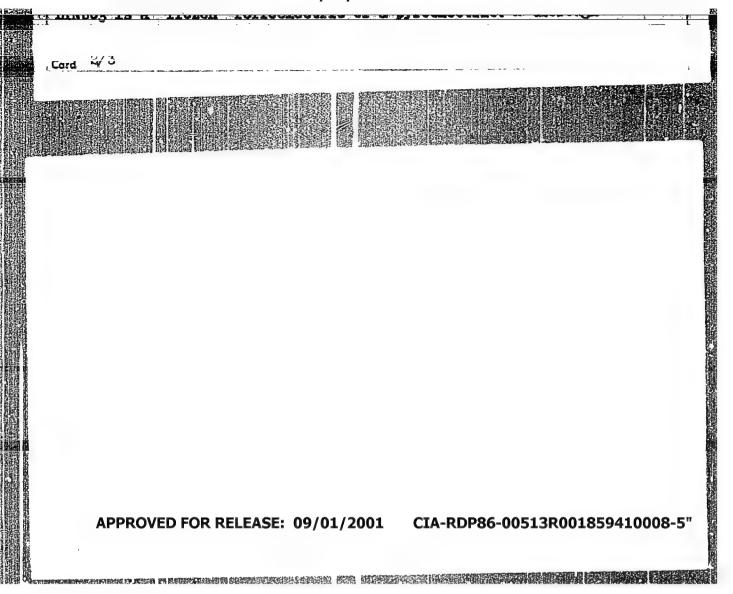


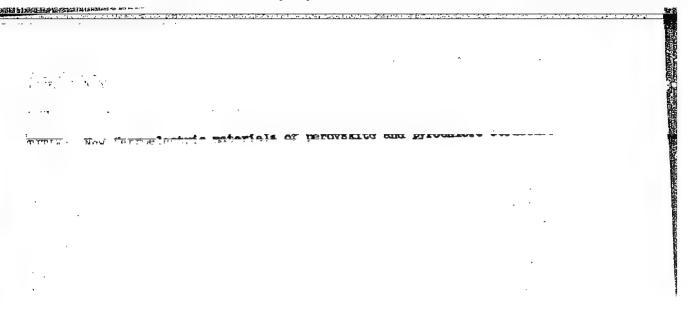


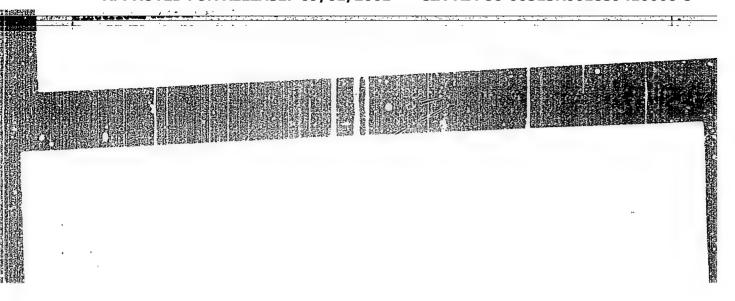


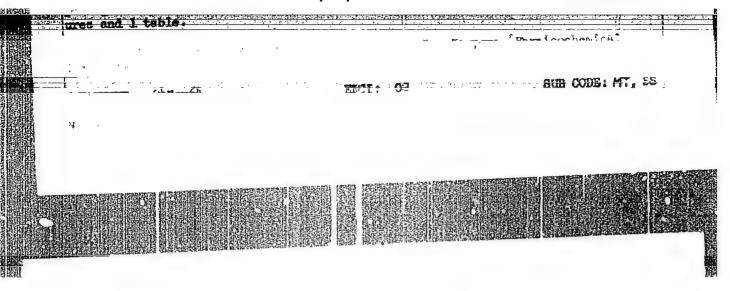


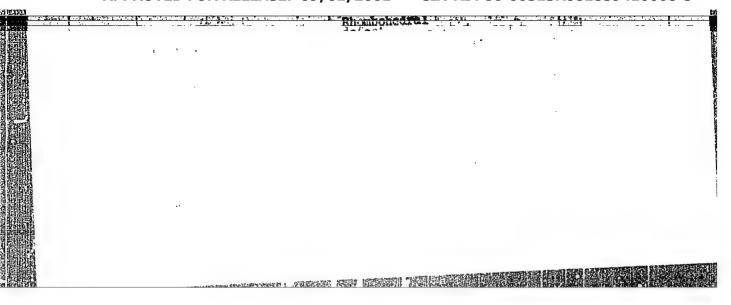


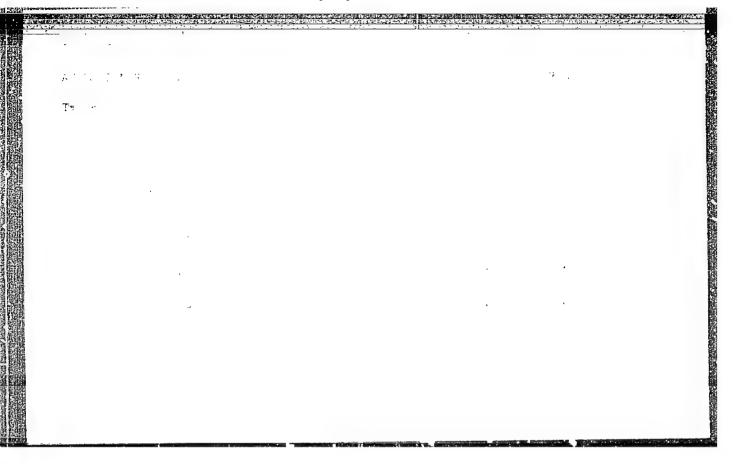


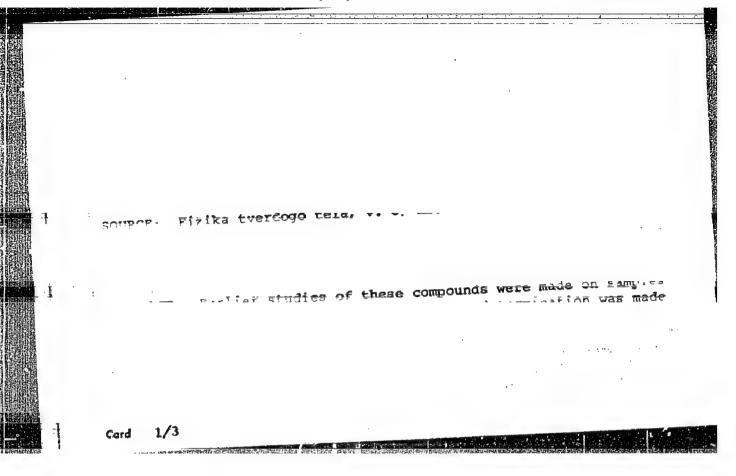


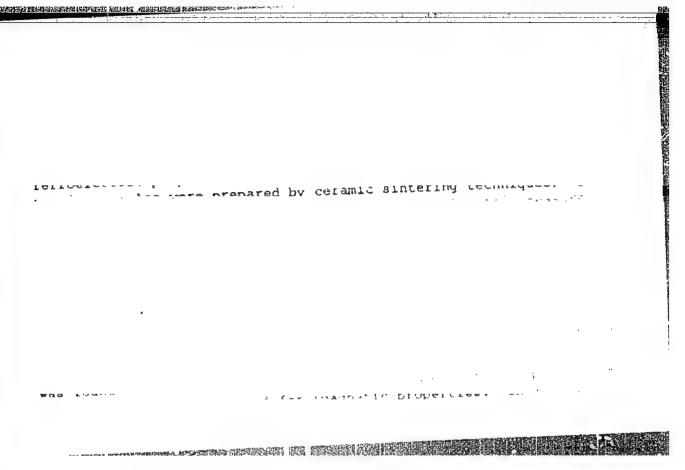


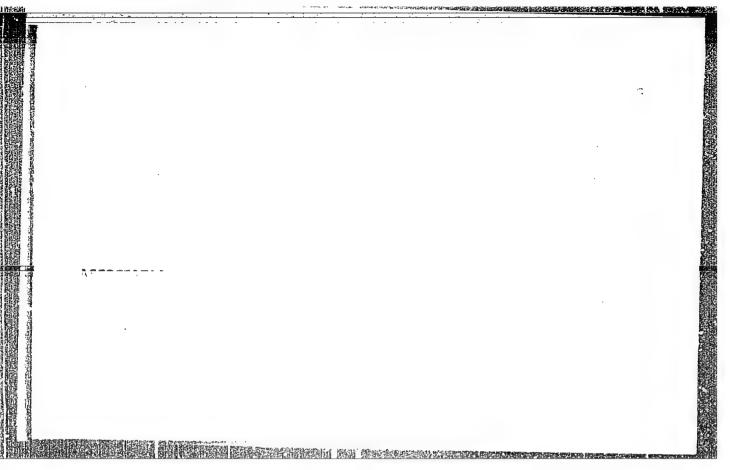












VENEVTSEV, Ya.N.; ROGINSKAYA, Yu.Ye.; VISKOV, A.S.; IVANOVA, V.7.;
TOMASHPOLISKIY, Yu.Ya.; SHVOMIEVA, L.I.; KAPYSHEV, A.G.;
TEVEROVSKIY, A. Yu.; ZHDANOV, G.S.

New lead-containing porovskite compounds of complex composition. Dokl. AN SSSR 158 no.1:86-38 S-0 *64 (MIRA 17:8)

1. Fiziko-khimicheskiy institut imeni L. Ya. Karpova. Prej-stavleno akademikom N.V. Belovym.

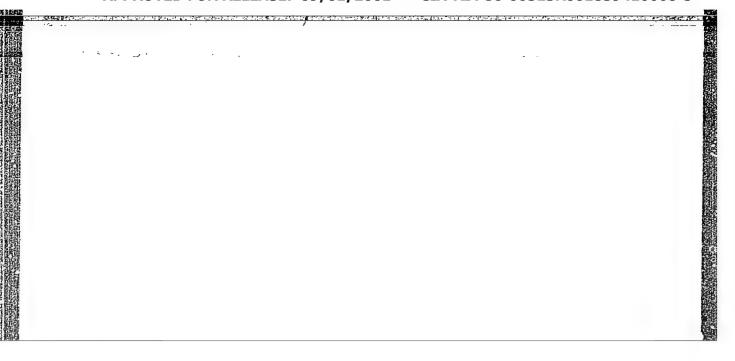
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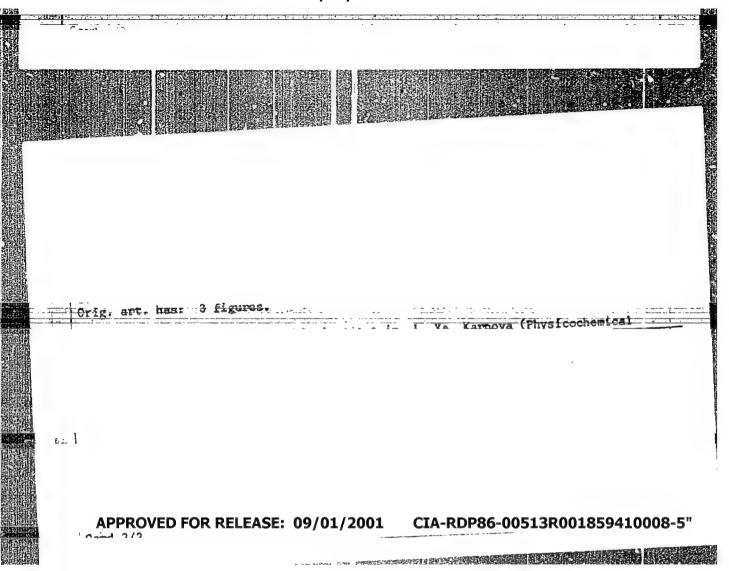
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VENGEROVSKIY, I.S., prof. (Tomsk)

Review of the book "Problems of pediatric surgery". (MIRA 17:11)

Review of the book "63.





VENEVISEV, Yu. N.; LYUBIMOV, V. N.; SOLOV'YEV, S. P.; ZHDAROV, G. S.

Calculation of internal electric fields and their gradients in perceptite compounds with specific dielectric properties. Izv. AN SSSR. Ser. fiz. 28 no. 4:630-635 Ap '64.

(MIRA 17:5)

MITROFAHOV, K.P.; VISKOV, A.S.; DRIKER, G.Ya.; PLOTNIKOVA, M.V.; FAM ZUI KHIYEN; VENEVTSEV, Yu.N.; SHPINEL', V.S.

Changes in the resonance absorption spectra of 23.8 Kev. gamma rays from Sn¹¹⁹ during phase transitions in the system BiFeO₃—Sr(Sn_{1/3} Mn_{2/3})O₃. Zhur. eksper. 1 teor. fiz. 46 (MIRA 17:2) no.1:383-386 Ja¹64.

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo universiteta i Fiziko-khimicheskiy institut imeni Karpova.

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TOMASHPOL'SKIY, Yu.Ya.; VENEVTSEV, Yu.N.; ZHDANOV, G.S.

Electron diffraction study of the crystalline structure of BiFeO3. Dokl. AN SSSR 153 no.6:1313-1314 D 163.

(MIRA 17:1)

1. Fiziko-khimicheskiy institut im. L.Ya. Karpova. Predstavleno akademikom N.V. Belovym.

SOLOVYEV, S. P.; LYUBIMOV, V. N.; VENEVTSEV, Yu. N.; ZHDANOV, G. S.

"The calculations of the internal electric fields and electric-field gradients in the perovskite-type compounds with special dielectric properties."

report submitted for 6th Gen Assembly, Intl Unionof Crystallography, Rome, 9 Sep 63.

ACCURACION DE LA CONTRACTOR DE LA CONTRA

Karpov Inst of Physical Chemistry, Moscow.

VENEVTSEV, Yu. N.; ZHDANOV, G. S.; ROGINSKAYA, Yu. Ye.; FEDULOV, S. A.; IVANOVA, V. V.

"Investigation of some solid solutions based on the ferroelectric-antiferromagnetic BiFeO 3."

report submitted for 6th Gen Assembly, Intl Union of Crystallography, Rome, 9 Sep 63.

Karpov Inst of Physical Chemistry, Moscow.

FEDULOV, S.A.; LADYZHINSKIY, P.B.; PYATIGORSKAYA, L.I.; VENEVTSEV, Yu.N.

Complete phase diagram of the system PbTiO3 - BiFeO3. Fiz. tver.

(MIRA 17:2)
tela 6 no.2:475-478 F '64.

1. Vsecoyuznyy nauchno-issledovatel'skiy institut khimicheskikh
reaktivov i osobo chistykh khimicheskikh veshchestv, Moskva.

SHAPIRO, Z.I.; FEDULOV, S.A.; VENEVISEV, Yu.N.

Curie temperature of the ferroelectric LiTaO3. Fiz. tver. tela 6 no.1:316-317 Ja '64. (MIRA 17:2)

1. Vsesoyuznyy nauchno-issledovateliskiy institut khimicheskikh reaktivov i osobo chistykh khimicheskikh veshchestv, Moskva.

ACCESSION NR: AP4012566 AUTHORS: Mitrofanov, K. P.; Viskov, A. S.; Driker, G. Ya.; Plots/0056/64/046/001/0383/0386 nikova, M. V.; Fam, Zui Khiyen; Venevtsev, Yu. N.; Shpinel', V. S. TITLE: Change in resonance absorption spectra of 23.8 keV gamma rays of Sn-119 during phase transitions in the system BiFeO3-Sr(Sn_{1/3}Mn_{2/3})0₃" SOURCE: Zhurnal eksper. i teoret. fiz., v. 46, no. 1, 1964, 383-386 TOPIC TAGS: resonance absorption, Mossbauer effect, recoilless resonance absorption, ferroelectric antiferromagnetic compound, ferroelectricity, ferro antiferromagnetism, group II stannate, resonance absorption maximum, resonance absorption jump, Mossbauer effect jump, magnetic hyperfine splitting ABSTRACT: This is a continuation of an earlier investigation by some of the authors (ZhETF v. 44, 2182, 1963) and is aimed at im-

ACCESSION NR: AP4012566

proving the earlier results and finding the reason for the abrupt change in the relative counting rate at the absorption maximum (ɛ). The material used has properties similar to that of the earlier investigation, and the addition of manganese made the samples practically single-phase and closer to equilibrium. The test procedure is briefly described. The results indicate that the jump in the value of the Mossbauer effect in solid solutions based on BiFeO₃ is the result of magnetic hyperfine splitting (but is not caused by change in the probability of the effect), and is related to an antiferromagnetic phase transition. This conclusion is supported by magnetic measurement results. Orig. art. has: 3 figures.

ASSOCIATION: Institut yadernoy fiziki Moskovskogo gosudarstvennogo universiteta (Nuclear Physics Institute, Moscow State University); Fiziko-khimicheskiy institut im. L. Ya. Karpova (Physicochemical Institute)

SUBMITTED: 27Sep63:

DATE ACQ: 26Feb64

ENCL: 02

Card 2/8/2

ACCESSION NO: AP4013507

S/0181/64/006/002/0475/0478

AUTHORS: Fedulov, S. A.; Lady*shinskiy, P. B.; Pyatigorskaya, L. I.; Veneytsev, Yu. N.

TITLE: Complete phase diagram of the system PbTiO3 BiFeO3

SOURCE: Fizika tverdogo tela, v. 6, no. 2, 1964, 475-478

TOPIC TAGS: phase diagram, PbTiO sub 3, BiFeO sub 3, piezoelectric, phase transition, Curie point, morphotropic phase transition, polarization, ferroelectric, ferromagnetic, Neel temperature, conductivity

ABSTRACT: Using x-ray investigations and electrical and magnetic measurements, the authors have constructed a complete phase diagram of the system PbTiO₃-BiFeO₃. This diagram is shown in Fig. 1 on the Enclosure. It is seen that in the region of the morphotropic phase transition the Curie point is very high (on the order of 700C), and it therefore seems suitable (in order to obtain high-temperature piezoelectric material) to introduce other material into the system to decrease the conductivity and to improve the conditions of polarization. The authors suggest, from this point of view, studies of the three-component systems PbTiO₃-BiFeO₃-PbZrO₃ and PbTiO₃-BiFeO₃-LaAlO₃. "The authors consider it their duty to express Card 1/4-7

ACCESSION NO: AP4013507

thanks to Yu. Ye. Roginskaya for her aid in the work." Orig. art. has: 5 figures.

ASSOCIATION: Vsesoyuzny*y nauchno-issledovatel'skiy institut khimicheskikh reaktivov i osobo chisty*kh khimicheskikh veshchestv; Moscow (All-Union Scientific Research Institute of Chemical Reagents and Extra Pure Chemical Substances)

SUBMITTED: 12Aug62

DATE ACQ: 03Mar64

ENCL: 01

SUB CODE: PH

NO REF SOV: 009

OTHER: 001

Card 2/3 2-

ACCESSION NR: APAO11779

5/0181/64/006/001/0316/0317

AUTHORS: Shapiro, Z. I.; Fedulov, S. A.; Venewtsev, Yu. N.

TITLE: Curie point of ferroelectric lithium tentalate

SCURCE: Fizika tverdogo tela, v. 6, no. 1, 1964, 316-317

TOPIC TAGS: reproelectric lithium tantalate, Curie point, dielectric property, lithium carbonate, tantalum pentoxide, lattice parameter, hexagonal exis, rhombohedral axis, dielectric permeability, bridge MPP-300, piezoelectric effect

ABSTRACT: An attempt was made to determine the dielectric properties in ceremic specimens of lithium tantelate across a broad temperature range. Experimental specimens were produced from lithium carbonate and tantalum pentoxide. They were purified by two repeated heatings (60 min each), one at 1100C and one at 1350C. X-ray analysis and subsequent calculations proved that the lattice parameters of

lithium tantalate were: on hexagonal axes -- $a_H = 5.153$ Å and $c_H = 13.775$ Å; on rhombohedral axes -- $a_{Rh} = 5.470$ Å and $\alpha_{Rh} = 56^{\circ}12^{\circ}$. Dielectric permeabilities

Card 1/2

ACCESSION NR: APAOL1779

were measured with a bridge MPP-300 at the frequency of 250 kilohertz. The curve of E = f(T) showed a sharp maximum at the temperature of about 665C. Dielectric permeability at room temperature was 70, at the maximum it reached 1850. Above the Curie point the change in the dielectric permeability was calculated from the Curie-Weiss law. The arccimens showed a weak piezoelectric effect. These results disprove the claim made by H. D. Megave (Acta Cryst., 7, 191, 1954; "Ferroelectricity in crystals," p. 103, London, 1957), to the effect that lithium tantalate forms simple pyroelectrical crystals. At the present time the authors are undertaking a study of properties exhibited by LiTaO₃ and LiNbO₃ and also of solid solutions based on these substances. Orig. art. has: 1 formula and 1 diagram.

ASSOCIATION: Vsesoyuzny*v nauchno-issledovatel*skiy institut khimicheskikh reaktivov i osobo chisty*kh khimicheskikh veshchestv, Moscow (All-Union Scientific Research Institute of Chemical Reactions and of Pure Chemical Materials)

SUBMITTED: 12Aug 63

DATE ACQ: 14Feb64

ENCL: 00

SUB CODE: CH. PH

NO REF SOV: OOO

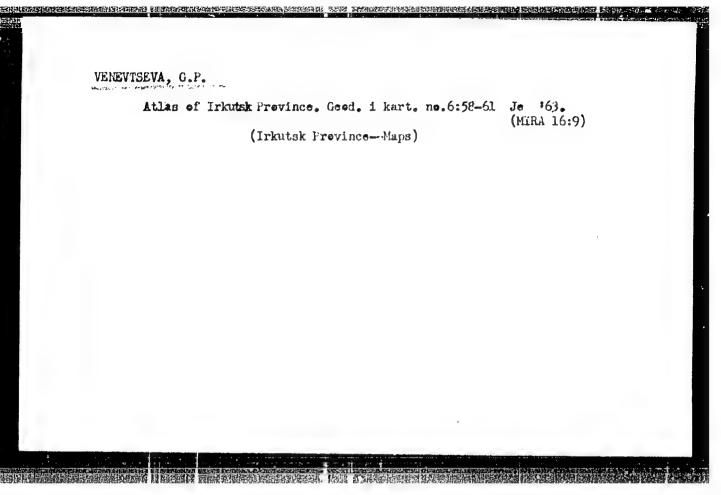
OTHER: 002

Cord 2/2

FET OV, S.A.; VENEVTSEV, Yu.N.

Transition region between the ferroelectric and paraelectric phases of the solid solutions (Pb, Ba) (Ti, Zr)03. Kristallografiia 8 no.3:454-456 My-Je 163. (MIRA 16:11)

1. Fiziko-khimicheskiy institut imeni L.Ya. Karpova.



AC	SOURCE CODE: UR/0070/66/011/005/0751/0755
AU OR TI	THOR: Tomashpol'skiy, Yu. Ya.; Venevtsev, Yu. N. 2: Physicochemical Institute im. L. Ya. Karpov (Fiziko-khimicheskiy institut) 3: Physicochemical Institute im. L. Ya. Karpov (Fiziko-khimicheskiy institut) 4: Physicochemical Institute im. L. Ya. Karpov (Fiziko-khimicheskiy institut) 5: The purpose of a system companet 6: TAGS: x ray diffraction, solid solution, ferroelectric material, ferromagnetic eterial, electric property, magnetic property, temperature dependence 7: The purpose of the investigation was to check on the feasibility of syntesizing compounds which exhibit simultaneously ferroelectric anf ferromagnetic properties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by the authors in earlier investigations (FTT v. 7, exties, which was demonstrated by sintering the extinction of the samples were determined by corresponding oxides. The properties and structure of the samples were determined by corresponding oxides. The properties and structure of the samples were determined by corresponding oxides. The properties and structure of the samples were determined by corresponding oxides. The properties and structure of the samples were determined by corresponding oxides. The properties and structure of the samples
1	UDC: 537.226

Phase diagrams of the two systems are presented. The results show that the compounds remain ferromagnetic in almost the entire range of concentrations, and that compounds remain ferromagnetic properties at 87 - 95 and 90 - 97 mol. respectively. I and II have ferroelectric properties at 87 - 95 and 90 - 97 mol. respectively. Although the ferroelectricity and the ferromagnetism combine only at low temperatures, Although the ferroelectricity and the ferromagnetism combine only at low temperatures, at it is assumed that the temperature range can be extended by suitably choosing the initial components. The authors thank V. P. Glushkov and A. I. Abramov for chemical

analysis. Orig. art. has: 5 figures.

SUB CODE: 20/ SUBM DATE: 12May65/ ORIG REF: 005/ OTH REF: 002

是**,我们就们是没有的**,我们就是我们的,我们就是这个人,我们就是我们的,我们就是我们的,我们就是这个人,我们就是这个人,我们就是这个人,我们就是这个人,我们就是

Cord 2/2

SEMENOV, A.I., otv.red.; FILIPPOV, Yu.V., prof., doktor tekhn.nauk, red.; BASHLAVIN, V.A., kand.tekha.nauk, red.; VOYNOVA, V.V., red.; GURARI, Ye.L., kand.ekonom.nauk, red.; GUREVICH, I.V., red.; ZHIV, I.S., red.; ZARUTSKAYA, I.P., red.; ZASLAVSKIY, I.I., red.; KOZLOV, F.M., red.; HIKISHOV, M.I., kand.geograf.nauk, red.; SADCHIKOV, S.F., red.; TIKHOMIROV, D.I., red.; TUTOCHKINA, V.A., red.; BALANTSEVA, I.A., red. kart; BOGDANOVA, L.A., red.kart; BOCHAROVA, I.L., red.kart; VENEVISEVA, G.P. red.kart; VOLKOVA, A.P., red.kart; GOSTEVA, N.A., red.kart; YEFIMOVA, G.N., red.kart; ZHIV, D.I., red.kart; KRAVCHENKO, A.V., red. kart; KUBRIKOVA, N.S., red.kart; KUZNETSOVA, N.A., red.kart; KURSAKOVA, I.V., red.kart; LOBZOVA, N.A., red.kart; MERTSALOVA, L.H., red.kart; MOSTMAN, S.L., red.kart; PANFILOVA, M.V., red.kart; SEMENOVA, V.D., red.kart; SMIRHOVA, T.H., red.kart; TERESHKOVA, V.S., red.kart; FEDOROVSKAYA, G.P., red.kart; FETISOVA, N.P., red.kart; FIL'GUS, Z.Kh., red.kart; SHAPIRO, Ye.M., red.kart; SHISHKIN, Ye.A., red.kart; YASHU-NICHKINA, Ye.G., red.kart. V razrabotke kart prinimali uchastiye: ALISOV, B.A., prof.: BERZINA, M.Ya.; VASILEVSKIY, L.I.; GAVRILOVA, S.A., kand.geograf.nauk; GINZBURG, G.A., kand.tekhn.nauk; DOBOSHINSKAYA, I.B.; YEVSTICHEYEVA, A.I.; LAVRENKO, Ye.M., prof.; LOZINOVA, V.H., kand. tekhn.nauk; MILANOVSKIY, Ye.Ye., kand.geologo-mineral.nauk; HIKHAYLOV, A.A., prof.; MYSHKIN, Ye.P.; PUZANGVA, V.F., kand.geograf.nauk; (Continued on next card)

APPROVED FOR RELEASE: 09/01/2001 CIA-RDP86-00513R001859410008-5"

SEMENOV, A.I.——(continued) Card 2.

ROZOV, N.N., prof.; SMIRNOV, D.I.; TARASOV, A.P.; TROFIMOVSKAYA.

Ye.A., kand.geograf.nauk; TUGCLESOV, D.A., kand.geologo-nineral.

nauk. ZININ, I.F., tekhn.red.

[Geographical atlas for secondary school teachers] Geograficheskii atlas; dlia uchitelei srednei shkoly. Izd.2. Moskva, Glav.upr. geodezii i kartografii MVD SSSR, 1959. 191 p. (MIRA 12:11)

1. Predstavitel' Nauchno-issledovatel'skogo instituta metodov obucheniya Akademii pedagogicheskikh nauk RSFSR (for Zaslavskiy). 2. Predstavitel' Upravleniya shkol Ministerstva prosvyashcheniya RSFSR (for Tutochkina). 3. Chleny-korrespondenty AN SSSR (for Lavrenko, Mikhaylov).

(Maps)

GRINEV, A.N.; VENEVTSEVA, N.K.; FRANCHUK, V.I.; TERENT'YEV, A.P.

Quinone.. Part 31: Synthesis of tetrahydro-1,4-endomethy-leneanthraquinones. Zhur.ob.khim. 30 no.6:1911-1914
Je '60. (MIRA 13:6)

Moskovskiy gosudarstvennyy universitet.
 (Anthraquinone)

GRINEV, A.N.; ZAYTSEV, I.A.; VENEVTSEVA, N.K.; TERENT'YEV, A.P.

Quinones. Part 32: Synthesis of substituted 2,5-bis(amino) -1.4-benzoquinones and 2-amino-,4-napthoquinones. Zhur.ob. khim. 30 no.6:1914-1918 Je '60. (MIRA 13:6)

大学的一个工作,这个工作,我们也没有这些证明的这种证明的证明,我们可以是一个工作,这个工作,这一个工作,这一个工作,我们是一个工作,我们们还是一个工作,我们们的证明,我们

1. Moskovskiy gosudarstvennyy universitet.
(Benzoquinone) (Maphthoquinone)

GRINEV, A.N.; VENEVTSEVA, N.K.

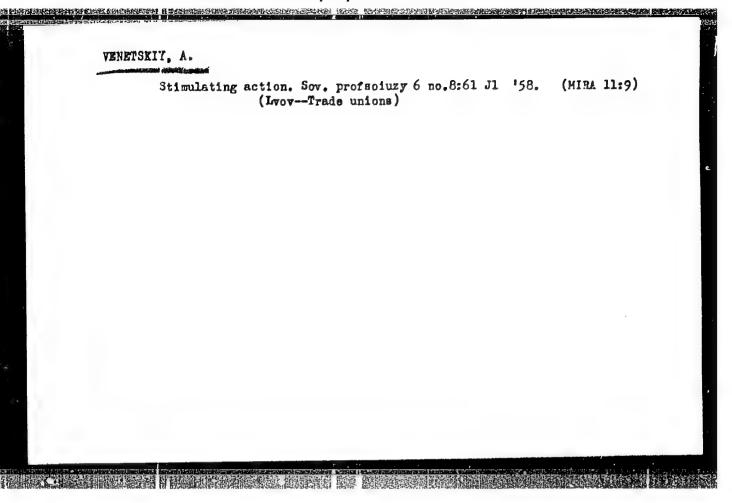
Aminomethyl and aminoethyl derivatives of 5-methoxybenzofuran. Zhur. ob. khim. 33 no.5:1436-1442 My 163. (MIRA 16:6)

1. Institut po izyskaniyu novykh antibiotikov AMN SSSR. (Benzofuran) (Amino group)

VENETSIANOV, Ye. A.; MOVEETOV, N. S.; BELIBASH, B. A.; SHIDLOVSKIY, M. F.; FEYVE,,
POLIKARPOV, P. 1.; BENG, A. Ya.; BLENHAN, A. A. and OBKENSKIY, V. Ye.;

"Tre Case for Explosion-proof Electrical Equipment in the Oil and Gas Industries."

report presented at the All Union Scientific and Technical Conference on the Electrical Equipment in Buildings and Outside Installations Liable to Explosions, 14-19 April 1958, Stalino (Energet, Byulleten', 1958, No. 7, pp 29 33).



DETORE, M.F.; VENETSKIY, V.L.

Optical properties of Ty -centers in ionic crystals. Fiz. shor.

(NIRA 11:8)

1. Institut fixiki AN USSR.

(Ionic crystals—Optical properties)

VENEV. M.

Myositis ossificans circumscripta neurotica. Khirurgiia, Sofia 11 no.1:84-88 1958.

l. Iz nevrokhirurgichnoto otdelenie pri Obshchoarmeiskata bolnitsa - Sofiia.

(MYOSITIS OSSIFICANS, circumscripts neurotica (Bul))

THE PROPERTY OF THE PROPERTY O

VENGLINSKIT, I.V. [Venhlins'kyi, I.V.]

New data on Spirialis from middle Miocene deposits of Trans
How data on Spirialis from middle Miocene deposits of Trans
Geol. shur. 18 no.5170-75 '58. (MIRA 12:1)

(Transcarpathia—Pteropoda, Tossil)

VEHBITSKAYA, T.N.; ZHDANOV, G.S.; VENEVISEV, Ku.N.; SOLOV'IEV, S.P.

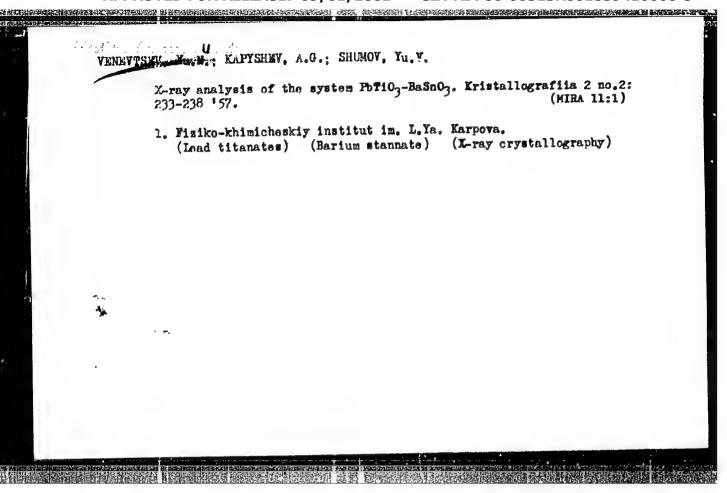
Hiectric testing and L-ray diffraction examination of the
Halio - Bazo system. Kristallografiia 3 no.2:186-196 '58.

(MIRA 11:6)

1. Fiziko-khimicheskiy institut im. L.Ia. Karpova i Kauchneissledovatel skiy institut Ministerstva radiotekhnicheskoy
promychlennosti.

(Barium titanate)

(Barium sirconate)



ZHDANOV, G.S.; SOLOV'YEV, S.P.: VENHVISHV, Yu.N.

,i.,

Structural coefficients of internal fields in seignetto-electric substances with perovskite-type structures. Kristallografia 2 no.5:639-645 \$57. (MIRA 11:1)

1. Fiziko-khimicheskiy institut im. L.Ya. Karpova. (Ferroelectric substances) (Grystal lattices)

USSR/Electricity - Dielectrics

G-2

Abs Jour

: Ref Zhur - Fizika, No 1, 1958, 1273

Author

: Venevtsev, Yu.M., Zhdanov, G.S.

Inst Title

: - : Crystal Chemistry of Ferroelectrics with the Perovskite

Structure.

Orig Pub

: Izv. AN SSSR, ser. fiz., 1957, 21, No 2, 275-285

Abstract

: A classification is made of ferroelectric and antiferroelectric substances of composition ABO; with a structure of the type of perovskite, in accordance with the type of the atomic displacements. The authors investigate also the connection between the structure and the character of the spontaneous polarization. It is shown that the ferroelectric or antiferroelectric properties of the crystal are due to the ion that has a sufficient geometric freedom of displacement within the confines of the crystalline cell. In the cells of the ferroelectrics

Card 1/3

USSR/Electricity - Lielectrics

Abs Jour : Ref Zhur - Fizika, No 1, 1958, 1273 G-2

When t > 1, it is cation B that is free within the confines of the cell, and when t < 1 it is cation A. For ferroelectrics one observes values of t that are both greater and less than 1, for antiferroelectrics t is always less than 1. An investigation of the atomic displacements has shown that cations A or B can shift along the second, third, and fourth order axis and cause respectively monoclinic, rhombohedral, and tetragonal deformation of the cells, while the cells of the antiferroelectrics display only a monoclinic deformation. An investigation of the character of the chemical bond in compounds ADO3 with a structure of the perovskite type has shown that the characteristic bond for these compounds is of the ionic type.

Bibliography, 46 titles.

Card 3/3



VEHENTSEV, YO. N

70-5-12/31

Zhdanov, G.S., Solov'yev, S.P. and Venevtsev, Yu.H.

The Structural Coefficients of the Internal Field in Ferro-AUTHORS: electrics with the Perovskite-type Structure. (Strukturnyye TITLE:

koeffitsiyenty vnutrennego polya v segnetoelektilkakh so

strukturoy tipa perovskita) Kristallografiya, 1957, Vol.2, No.5, pp. 639-645 (USSR)

USTRACT: Data published in the literature for the values of the PERIODICAL: structural coefficients of the internal field in perovskitetype crystals are critically examined. Relations are set up between the idealised cubic perovskite and the cases in which there are dipoles in the [OOI], [OII] or [III] directions. The structural coefficients are calculated for the tetragonal cell of PhTiO, at room temperature taking account of the ionic displacements.

The field at the i-th ion is:

$$E_{i} = E + \sum_{k=1}^{m} \left(\frac{4\pi r}{3} + C_{ik} \right) p_{k}$$

where m is the number of sub-lattices each consisting of the ions of the k-th sort, E is the external field, Pk is the

Card1/4

共和国最低<mark>的制度。1965年的</mark>特别的特别的基础的1660年的基础的1960年的

70-5-12/31

The Structural Coefficients of the Internal Field in Ferroelectrics with the Perovskite-type Structure.

dipole moment of the k-type of ions, C_{ik} are the structural coefficients. In the case where the dipoles are in the z-direction:

 $c_{ik} = \sum_{j} \frac{2z_{jk}^{2} - x_{jk}^{2} - y_{jk}^{2}}{\left(\frac{z_{jk}^{2} + y_{jk}^{2} + z_{jk}^{2}\right)5/2}$

where x_{jk} , y_{jk} , z_{jk} are the co-ordinates of the j-th dipole of the k-th sort relative to a dipole of the i-th sort and summation is over each dipole of the k-th sort. For the cubic cell, the C_{ik} can be expressed in terms of two quantities P and Q. The values which various authors find for these values are compared, the best values being P = -15.04102/V and Q = 4.33387/V as found by values being P = -15.04102/V and Q = 4.33387/V as found by NcKeehan (Phys. Rev. 43, 913, 1933 and 72, 78, 1947).

Card2/4

70-5-12/31

The Structural Coefficients of the Internal Field in Ferroelectrics with the Perovskite-type Structure.

Where the distortions of the cubic cell are small (1%) the C_{ik} are different from the C_{ik} only by 2-3%. Even for PoTiO₃ where the distortions are large, these do not differ by more than 20%. The actual values of the coefficients for truragonal PbTiO₃ where c/a = 1.064 are calculated and tabulated with those of McKeehan (cubic, by Ewald's method) and of Hagendorn (BaTiO₃ with c/a = 1.010) (Zeit.f.Physik, 133, 394-421, 1952). There are 1 figure, 3 tables and 14 references, 2 of which are Slavic.

ASSOCIATION: Karpov Physico -chemical Institute (Fiziko-khimicheskiy Institut im. L.Ya. Karpova)

The Structural Coefficients of the Internal Field in Ferroelectrics with the Perovskite-type Structure.

SUBMITTED: May 31, 1957.

AVAILABLE: Library of Congress

Card 4/4

VENEVISEYA, G.P. Using the dotted method to map branches of agriculture (practice followed in making maps for the Agricultural Atlas of the U.S.S.R.). Geod,1 kart, no.4357-61 Ap '62. (MIRA 15:12) (Agriculture—Maps) (Cartography)

VENEVISEVA, G.P. Using the dotted method to map branches of agriculture (practice followed in making maps for the Agricultural Atlas of the U.S.S.R.). Geod.i kart. no.4457-61 Ap '62. (MIRA 15:12) (Agriculture—Maps) (Cartography)

5.3610

77376

AUTHORS:

SOV/79-30-1-37/78

TITLE:

Grinyev, A. N., Venevtseva, N. K., Terent'yev, A. P.

Investigation in the Field of p-Qalnones. XXIX.

Azo Coupling of 2,5-bis (Dimethylamino)-1,4-benzoquinone and 2-(Dimethylamino)-1,4-napthoquinone

PERIODICAL:

Zhuenal obshchey khimii, 1960, Vol 30, Nr 1, pp 183-185 (USSR)

ABSTRACT:

Azo coupling of quinones containin; other electrodonor substituents was studied. 2,5-bis-(Dimethylamino)-

1,4-benzoquinone was coupled with some diazonium salts, and 2,5-bis-(dimethylamino)-3,6-bis-(p-nitrophenylazo)and 2,5-bis-(dimethylamino)-5,0-bis-(p-nitrophenylazo)-1,4-benzoquinone (III), 2,5-bis(dimethylamino)-3,6-bis-(o-nitrophenylazo)-1,4-benzoquinone (IV), and 1,4-benzoquinone (V) were obtained, in high yield.

Card 1/3

XXXIX.

77376 SOV/79-30-1-37/78

$$Ar = N = N$$

$$(CH_3)_2 N$$

$$(CH_3)_2 N = N = N$$

$$(V) Ar = V$$

$$(V) Ar = V$$

$$(V) Ar = V$$

Diazonium salts were reacted with 2-dimethylamino-1,4-naphthoquinone, and instead of the expected products of azo coupling of 2-dimethylamino-1,4-naphthoquinone, the derivatives of 2-hydroxy-1,4-naphthoquinone were obtained (2-hydroxy-3-(p-nitrophenylazo)-1,4-naphthoquinone (VI) and 2-hydroxy-3-(o-nitrophenylazo)-1,4-naphthoquinone (VII)).

Card 2/3

Investigation in the Field of p-Quinones. XXIX.

77376 \$07/79-30-1-37/78

III was obtained in 81.3% yield; IV, 38.6%; V, 65.9%; VI, 88.6%; VII, 77.1%. There is 1 table; and 11 references, 6 German, 3 U.S., 1 Italian, 1 U.K. The Chem. Soc., 56, 2478 (1934); L. Fieser, 1bid, 70, 3203 (1948); R. Baltzly, E. Lorz, 1bid, 70, 861 (1948); R. Plimpton, J. Chem. Soc., 37, 642 (1880).

ASSOCIATION:

Moscow State University (Moskovskiy gosudarstvennyy universitet)

SUBMITTED:

January 5, 1959

Card 3/3

和智慧和

GRINEV, A.N.; VENEVTSEVA, N.K.

Mannich bases in the series of derivatives of 5-hydroxybenzofuran. Zhur.ob.khim. 33 no.3:820-824 Mr '63. (MIRA 1613)

 Institut po izyskaniyu novykh antibiotikov AMN SSSR. (Mannich bases) (Benzofuran)

GRINEY, A.N.; VENEYSEVA, N.K.; TERENT'IEV, A.P.

Quinones, Part 14: The reaction of M -benzoquinone with M -nitroand M -bromobenzoylacetic esters. Zhur. ob. khim. 26 no.10:29332934 0 '56. (MIRA 11:3)

1. Moskovskiy Gosudarstvennyy universitet.
(Benzoquinone) (Acetic acid)

GRINAY, A.N.; VENETSEYA, N.K.; TERENTYZY, A.P.

Research in the field of quinones. Part 17: Reaction of 2,3dichloroquinone with benzoyl acetic ester. Zhur. ob. khtm. fr
no.h:1090-1091 Ap 157.

1. Moskovskiy gosudarstvennyy universitet.

(Quinone)

507/79-28-7-26/64

AUTHORS:

Grinev, A. N., Venevtseva, N. K., Werent yev, A. P.

TITLE:

The Alyklation of the Substituted 5-Oxybenzofurfuranes. The Synthesis of the New Plant Growth Stimulators (Alkilirovaniye zameshchennykh 5-oksibenzofuranov; eintez novykh stimulyatorov

PERIODICAL:

Zhurnal obshchey khimii, 1958, Vol. 28, Nr 7, FI. 1850-1853 rosta rasteniy)

(USSR)

ABSTRACT:

The authors succeeded in elaborating a method of the synthesis of the substituted 5-oxybenzofurfuranes by the condensation of p-quinone with the esters of the 2-keto scide (Refs 1 - 5), which practically made accessible a series of 5-oxybenzofurfurane derivatives. As a proof for the structure of the obtained compounds papers already existed on the methylation of some of these compounds with dimethyl sulfate, with either derivatives of the 5-methoxybenzofurfurane-3-carboxylic acids (Refs 3, 5), or their esters having been obtained, depending on the conditions prevailing (Ref 6). The compounds (I), (II), (III), (IV) and (V) were synthetized by the authors by the conversion of the 5-oxybenzofurfurane with dimethylsulfate

Card 1/2

The Alkylation of the Substituted 5-Oxybenzofurfurance. The Synthesis of the New Plant Growth Stimulators

in alkaline medium in the presence of dioxane. In the hydrolysis of the esters (I), (III), (V) and (VI) with alcoholate the furfuranes (VII), (VIII). (IX) and (X) were obtained. The acetic acids (XI) and (XII) were obtained as final products of the alkylation of the already earlier (Ref 1) synthetized oxy acids of the benzofurfurane series with chloroacetic acid. The potassium salts of the acids (VII) - (XII) proved to be highly effective plant growth stimulators we was shown by the evidence obtained by N. A. Bazilevskaya in the M. G. U. Botanical Garden. The results of the biological experiments with these products will be published later. There are 6 references, 6 of which are Soviet.

ASSOCIATION:

Moskovskiy gosudarstvennyy universitet (Moscow State University)

SUBMITTED:

June 18, 1957

Card 2/2

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50V/73-28-7-27/64

AUTHORS:

Grinev, A. H., Zaytsev, I. A., Veneytseva, H. K.,

Terent yev, A. P.

TITLE:

A New Method for the Synthesis of Substituted Benzofurfuraneand Indole From Esters of the Benzofurfurane- and Indole-2-Carboxylic Acids (Novyy metod polucheniya zameshchennykh benzofuranov i indolov iz efirov benzofuran-i indol-β-kar-

bonovykh kislot)

PERIODICAL:

Zhurnal obshchey khimii, 1958, Vol. 28, Nr 7, pp. 1853-1855

(USSR)

ABSTRACT:

In many cases the esters of the substituted benzofurfuraneand indole-3-carboxylic acids were bitter accessible than
the benzofurfuranes and indoles as such. The carboxyl group
in these compounds is closely attached to them so that their
in these compounds strict conditions which lead to a great
cleavage demands strict conditions which lead to a great
loss in substance (Refs 1, 2). In the experiments carried
out to remove the carboxyl group from such and similar compounds the authors either met with difficulties, or the
yields were too small (Refs 3 - 6). The synthesis of the
N-alkyl- and N-aryl substituted indoles with a free β-posi-

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507,79-28-7-27/64

A New Method of the Synthesis of Substituted Benzofurfurane- and Indole From Esters of the Benzofurfurane- and Indole-β-Carboxylic Acids

tion could not be realized in these experiments at all. The authors by means of some examples suggest a convenient method for the cleavage of the esters of the benzofurfurane—and indole-β-cmboxylic acids by their heating with equimolecular quantities of sulfuric or phosphoric acid in glacial acetic acid solution. The reaction takes place according to the mentioned scheme. Thus the authors by the action of sulfuric acid on the corresponding esters obtained the furfuranes (I), (II), (III) and the indole (IV), and by the action of phosphoric acid the indoles (V) and (VI). The 2-phenyl-j-carboxy-6,7-dichlorobenzofurfurane-5-oxyacetic acid was also subjected to the cleavage of sulfuric acid, with the compound (VII) having been obtained. There are 9 references, 6 of which are Soviet.

ASSOCIATION:

Moskovskiy gosudarstvennyy universitet

(Moscow State University)

SUBMITTED:

June 18, 1957

Card 2/3

Furan derivatives—Synthesis
 Substitution reactions

3. Plants-Growth 4. Growth substances--Synthesis

A New Method for the Synthesis of Substituted Benzofurfurane- and Indole From Esters of the Benzofurfurane- and Indole-6-Carboxylic Acids

1. Furan derivatives—Synthesis 2. Indoles—Synthesis 3. Carboxylic acid esters—Chemical reactions 4. Substitution reactions 5. Carboxyl radicals

--Chemical effects

Card 3/5

297/79-28-7-28/64

.uTHORS:

Grinev, A. N., Venevtseva, N. K., Terent'yev, A. P.

TITLE:

Investigations in the Field of Quinones (Iseledoveniya v oblasti khinonov) (XIII. The Investigation of the Condensation of p-Benzoquinone and 2,3-Dichloroquinone With Acetic Ester and Its Analogs (XXIII. Izucheniya kondensatzii p-benzo-khinona i 2,3-dikhlorkhinona s atsetoukenenym efirom i yego

enalogumi)

和我们**的自己和解释的对比**解现象。对比例将是是对 自由显然是对此的影影的,我就是不是是一个人的一个人,就是一个人的一个人,就是一个人的一个人,就是一个人的一个人

PERIODICAL:

Zhurnal obshchey khimii, 1958, Vol. 28, Nr 7, pp. 1856-1864

(USSR)

ABSTRACT:

In the present paper the authors carried out the condensation of the p-benzoquinone with acetic ester at lower temperatures than was the case in their earlier papers (Refs 1 - 5); the than was the case in their earlier obtained. At 80-85 and results deviated from those earlier obtained. At 80-85 and at a low concentration of p-benzoquinone in the reaction mixture mainly (at low concentrations even exclusively) a benzofurfurane derivative (formula I) is obtained, at 41-45 at low concentrations the esters (II) and (III) are formed. When the reaction takes place at 38 only the substituted

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307/79-28-7-28/64

Investigations in the Field of Quinones. XXIII. The Investigation of the Condensation of p-Benzoquinone and 2,3-Dichloroquinone With Acetic Ester and Its Analogs

benzofurfurylacetic ester (II) is obtained without admixture of (III). The substituted ester (II) on boiling its solutions in alcohol and other solvents easily converts to (III). When the reaction is carried out at low temperatures it is possible to separate also a product with a melting point of 206.5 - 2070 besides (II) from the reaction products when the relative concentration of p-benzoquinone is increased; according to the analysis and the qualitative reactions carried out this product must be given the formula (IV). From the experiments carried out it may be concluded that the one or the other direction of the condensation of the quinones with esters of the f-keto acids, which may be easily controlled by the comparison of the yields of the derivatives of benzofurfurane and benzodifurfurane, does not only depend on the concentration of quinone in the reaction mixture but also to a considerable degree on temperature. Besides these factors mentioned also the nature of the reagent acting in the reaction with the quinones influences the yield of the derivatives of benzofurfurane and benzodifurfurane. There are 8 references,

Card 2/3

117/79-28-7-38/64

Investigations in the Field of Minones. KMIII. The Investigation of the Condensation of p-Benzoquinone and 2,5-bicklorocuinone litt Acetic Ester and Ite snalogs

7 of which are deviet.

ASSOCIATION: Moskovskiy gosudaratvennyy universitet

(Moscow State University)

SUBMITTED:

June 21, 1957

1. Benzoquinones--Chemical reactions 2. Chloroquinone--Chemical 3. Condensation reactions 4. Acetic acid esters-Chemical reactions

reactions

Card 3/3